Ser. No. 10/597,954 Response to Office Action of 3 Aug 2009

Atty Docket 114116.00032

AMENDMENTS TO THE CLAIMS

Listing of Claims:

1. (currently amended) An apparatus for separating and purifying nucleic acids

comprising an integral monolith structure, wherein macro-pores continuously extending

from one end of the monolith structure to the other end and corresponding to the sizes of

nucleic acids are provided and configured so that nucleic acids corresponding to the macro-pores can be retained respectively by allowing a solution containing nucleic acids

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to be separated to pass therethrough, wherein the diameter range of the micropores

macro-pores is selected according to the size of the nucleic acid to be purified and wherein the size range is selected from the group consisting of diameters of about 10

nanometers (nm) to about 100 nm, diameters of about 100 nm to about 1 micrometers

anomore (min) to access 100 min, claimeters of access 100 min to access 1 micromoters

(µm), diameters of about 1 µm to about 10 µm, and diameters of about 10 µm to about

 $100\;\mu\text{m},$ and further wherein the monolith structure is capable of adsorbing nucleic acids

in the presence of potassium ions and is capable of releasing nucleic acids in an

essentially salt-free solution.

2. (previously presented) The apparatus for separating and purifying nucleic acids

according to claim 1, wherein the monolith structure employs a glass, a silica or a hybrid material containing an organic material and a glass or a silica, which is a porous body

having macro-pores penetrating from an upper surface to a lower surface.

3. (previously presented)The apparatus for separating and purifying nucleic acids

according to claim 2, wherein the porous body of the monolith structure has micro-pores

in the macro-pores.

4. (currently amended) The apparatus for separating and purifying nucleic acids

according to claim 3, wherein the porous body of the monolith structure has a micro-pore size of greater than zero and less than or equal to 100 nm.

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(previously presented) The apparatus for separating and purifying nucleic acids according to claim 1, wherein a disc formed with the monolith structure is placed in a column tube to form a monolith solid phase column.

6. (previously presented)The apparatus for separating and purifying nucleic acids according to claim 1, wherein the apparatus employs a monolith solid phase column formed by detachably attaching a base formed with the monolith structure to a cylindrical body having the top and the bottom opened.

(previously presented) The apparatus for separating and purifying nucleic acids
according to claim 1, characterized in that the porous body of the monolith structure has
micro-pores in the macro-pores.

 (previously presented) The apparatus for separating and purifying nucleic acids according to claim 1, wherein the porous body of the monolith structure additionally has a micro-pore size of 100 nm or less.

 (previously presented) The apparatus for separating and purifying nucleic acids according to claim 2, wherein the porous body of the monolith structure additionally has a micro-pore size of 100 nm or less.

10. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 2, wherein a disc formed with the monolith structure is placed in a column tube to form a monolith solid phase column.

11. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 3, wherein a disc formed with the monolith structure is placed in a column tube to form a monolith solid phase column.

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12. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 4, wherein a disc formed with the monolith structure is placed in a column tube to form a monolith solid phase column.

13. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 2, wherein the apparatus employs a monolith solid phase column formed by detachably attaching a base formed with the monolith structure to a cylindrical body having the top and the bottom opened.

14. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 3, wherein the apparatus employs a monolith solid phase column formed by detachably attaching a base formed with the monolith structure to a cylindrical body having the top and the bottom opened.

15. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 4, wherein the apparatus employs a monolith solid phase column formed by detachably attaching a base formed with the monolith structure to a cylindrical body having the top and the bottom opened.

16. (previously presented) The apparatus for separating and purifying nucleic acids according to claim 5, wherein the apparatus employs a monolith solid phase column formed by detachably attaching a base formed with the monolith structure to a cylindrical body having the top and the bottom opened.

17. (withdrawn, currently amended) A method for separating and purifying nucleic acids comprising a step of using an integral monolith structure, wherein macro-pores continuously extending from one end of the monolith structure to the other end and corresponding to the sizes of nucleic acids are provided and configured so that nucleic acids corresponding to the macro-pores can be retained respectively by allowing a solution containing nucleic acids to be separated to pass therethrough, wherein the

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macro-pores are selected according to the size of the nucleic acid to be purified, such that

macropores having a diameter of about 10 nanometers (nm) to about 100 nm are present

for separating and purifying nucleic acids of about 35 base pairs (bp) to about 300 bp, macro-pores having a diameter of about 100 nm to about 1 micrometers (µm) are present

for separating and purifying nucleic acids with about 300 bp to about 3 kilobase pairs

(Kbp), macro-pores having a diameter of about 1 μm to about 10 μm are present for

separating and purifying nucleic acids with about 3 Kbp to about 30 Kbp, and macropores having a diameter of about 10 µm to about 100 µm are present for separating and

purifying nucleic acids with about 30 Kbp to about 300 Kbp, and further wherein the

monolith structure is capable of adsorbing nucleic acids in the presence of potassium ions

and is capable of releasing nucleic acids in an essentially salt free solution.

18. (withdrawn) The method for separating and purifying nucleic acids according to

claim 17, wherein the monolith structure employs a glass, a silica or a hybrid material containing an organic material and a glass or a silica, which is a porous body having

macro-pores (through-pores) penetrating from an upper surface to a lower surface.

19. (withdrawn) The method for separating and purifying nucleic acids according to

claim 17, wherein the porous body of the monolith structure has micro-pores in the

macro-pores.

20. (withdrawn) The method for separating and purifying nucleic acids according to

claim 18, wherein the porous body of the monolith structure has micro-pores in the

macro-pores.

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